



□ Drug dispensed from a pharmacy (pharmacy benefit)

sunflower health plan. UnitedHealthcare'

# Kansas Medical Assistance Program

PA Phone 800-933-6593 PA Fax 800-913-2229

CHECK ONE:

### Aetna Better Health of KS

PA Pharmacy Phone 855-221-5656 PA Pharmacy Fax 844-807-8453 PA Medical Phone 855-221-5656 PA Medical Fax 855-225-4102

#### Sunflower

PA Pharmacy Phone 877-397-9526 PA Pharmacy Fax 866-399-0929 PA Medical Phone 877-644-4623 PA Medical Fax 888-453-4756

#### UnitedHealthcare

PA Pharmacy Phone 800-310-6826 PA Pharmacy Fax 866-940-7328 PA Medical Phone 866-604-3267 PA Medical Fax 866-943-6474

## **Asthma Agents PRIOR AUTHORIZATION FORM**

Complete form in its entirety and fax to the appropriate plan's PA department. For questions, please call the pharmacy helpdesk specific to the member's plan.

☐ Drug administered in an office or outpatient setting (medical benefit)						
MEMBER INFORMATION						
Name:		Medica	aid ID:			
Date of Birth:		Gende	Gender:			
PRESCRIBER INFORMATION	N					
Name:		Medica	aid ID:			
NPI:		Phone:		Fax:		
Address:		City, St	ate, Zip Code:			
The following medications require Prior Authorization (PA). Medications requiring PA may have to meet clinical and Non-Preferred PA criteria before the claim may be considered for payment.  Please provide the required data for the specific drug being requested. Below is a list of links you may find helpful in determining the required information:  Clinical PA criteria: <a href="http://www.kdheks.gov/hcf/pharmacy/pa_criteria.htm">http://www.kdheks.gov/hcf/pharmacy/ga_criteria.htm</a> KS Preferred Drug List (PDL): <a href="http://www.kdheks.gov/hcf/pharmacy/download/PDLList.pdf">http://www.kdheks.gov/hcf/pharmacy/download/PDLList.pdf</a> Non-Preferred, PA Required PDL criteria: <a href="http://www.kdheks.gov/hcf/pharmacy/download/NonPreferred PA Criteria for PDL Drugs.pdf">https://www.kdheks.gov/hcf/pharmacy/download/NonPreferred PA Criteria for PDL Drugs.pdf</a> KS NDC lookup tool: <a href="https://www.kmap-state-ks.us/Provider/PRICING/NDCSearch.asp">https://www.kmap-state-ks.us/Provider/PRICING/NDCSearch.asp</a> Note: Any area not filled out will be considered not applicable to this PA & may affect the outcome of this request.  Instructions to complete this form:  Complete the Member/Prescriber Information portion above and Section I for ALL requests.  Complete Section II if this request is also for a Non-Preferred PDL drug.  Complete Section III for all clinical information required.  Complete the section above and Section IV only, if the requested medication is a renewal.  Prescriber - Sign and date the form prior to submission.						
SECTION I: MEDICATION I						
Select the appropriate med Benralizumab (Faserna® Dupilumab (Dupixet ®) Mepolizumab (Nucala®) Omalizumab (Xolair®) Reslizumab (Cinqair®)	* *	est:				
NDC/HCPCS (J Code)	<u>Strength</u>	Dosage Form	<u>Quantity</u>	<u>Directions for Use</u>		
Indication/Diagnosis:						
Is the requested medication but Indication:		* *	□ YES □ NO			
Patient's weight:	. □ lbs. □ kg					

Providers: You are required to return, destroy or further protect any PHI received on this document pertaining to members whom you are not currently treating. Providers are required to immediately destroy any such PHI or safeguard the PHI for as long it is retained. In no event are you permitted to use or re-disclose such PHI.

PATIENT NAME:	MED	ICAID ID:				
SECTION II: NON-PREFERRED MED	NCATION					
Is the medication requested a non-preferred medication on the Kansas Medicaid preferred drug list (PDL)?  KS Preferred Drug List (PDL): <a href="http://www.kdheks.gov/hcf/pharmacy/download/PDLList.pdf">http://www.kdheks.gov/hcf/pharmacy/download/PDLList.pdf</a>						
	□ YES □ NO − Proceed to Section III					
If <u>YES</u> : Does the patient labeling as specified in t	have a documented clinical rationale for using a no he Non-preferred PDL PA criteria?	on-preferred medication that is supported by the product				
•	nred PDL criteria: <u>http://www.koneks.gov/nci/pharma</u> NO	cy/download/NonPreierred_PA_Criteria_lor_PDL_Drugs.pdi				
	ntation of clinical rationale to support the use of t	the requested non-preferred medication.				
SECTION III: CLINICAL INFORMATI	ON					
1. Is this a new or renewal requ						
□ New □ Renewal – Proceed						
	<ul><li>2. Please document the prescribing physician's specialty.</li><li>□ Pulmonologist □ Allergist □ Immunologist □ Other</li></ul>					
· <del></del>	cribing provider consulted with one of the provide ase document the provider's name, specialty and d					
Provider name	e: Specialty:	Date of Consult:				
□ NO						
<ol> <li>Please list all medications the patient is taking or has previously tried and failed for treatment of this diagnosis.</li> <li>*Specify medication name, Action Taken (continue medication, discontinue medication due to inadequate response, contraindication, intolerance) and dates of previous medication trial.</li> </ol>						
Medication name	Action Taken	<u>Dates of trial</u>				
4. Please list all medications the patient will use in combination with the medication requested for the treatment of this diagnosis.  Medication name(s):						
5. Does the prescriber attest that	at the patient is not currently on another biologic o	or janus kinase (JAK) inhibitor?				
6. Please provide the baseline F	EV <sub>1</sub> value (include units if applicable)					
7. Please provide the following i	7. Please provide the following information:					
<ul> <li>For all agents, number of Exacerbation is defined to the Control of the Control of</li></ul>	dates)					

Providers: You are required to return, destroy or further protect any PHI received on this document pertaining to members whom you are not currently treating. Providers are required to immediately destroy any such PHI or safeguard the PHI for as long it is retained. In no event are you permitted to use or re-disclose such PHI.

Revised 7/2019 Page 2 of 3

PATIENT NAME:	MEDICAID ID:	
SECTION III: CLINIC	CAL INFORMATION (continued)	
o F	malizumab, please provide one of the following: Perennial aeroallergen skin test result In vitro reactivity to a perennial aeroallergen	
0 (	upilumab, please provide one of the following: Oral Corticosteroid dose for corticosteroid-dependent Asthma Blood eosinophil count	
	enralizumab, Mepolizumab, and Reslizumab, please provide the following: Blood eosinophil count	
SECTION IV: RENE	WAL	
•	rescriber attest that the patient has received clinical benefit from continuous t  NO	reatment with the requested medication?
2. Please prov	vide one of the following:	
0	Most recent FEV <sub>1</sub> Value (Include Units if Applicable):	Date:
	Effect on Exacerbations in the past 12 months (with dates) Describe changes in exacerbation(s) compared to baseline:	
3. Please prov	vide the patient's current dose:	
4. Does the p	rescriber attest that the patient is not currently on another biologic or Janus k	inase (JAK) inhibitor?
□ YES □	□ NO	
PRESCRIBER SIGNA	ATURE	
☐ I have completed	d all applicable boxes and attached any required documentation for revie	w, in addition to signing and dating this form.
Prescriber or author	prized signature Date	
are appropriate for a pa	efits is not the practice of medicine or the substitute for the independent medical judgment of a treating pf atient. Please refer to the applicable plan for the detailed information regarding benefits, conditions, limite information provided is true, accurate, and complete and the requested services are medically indicated a Note: Payment is subject to member eligibility. Authorization does not guar	tions, and exclusions. The submitting provider certifies that the nd necessary to the health of the patient.

Providers: You are required to return, destroy or further protect any PHI received on this document pertaining to members whom you are not currently treating. Providers are required to immediately destroy any such PHI or safeguard the PHI for as long it is retained. In no event are you permitted to use or re-disclose such PHI.

Revised 7/2019 Page 3 of 3